PATENT COOPERATION TREATY

PCT/IL2004/000181

		From the INTERNATIONAL BUREAU		
PCT		То:		
NOTIFICATION CONCERNING TRANSMITTAL OF COPY OF INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (CHAPTER I OF THE PATENT COOPERATION TREATY) (PCT Rule 44bis.1(c)) Date of mailing (<i>day/month/year</i>)		G. E. EHRLICH (199 11 Menachem Begin 52 521 Ramat-Gan ISRAËL	5) LTRECEIVED 17 APR 2006 FILE No. <u>27558</u> G.E. EHRLICH (1995) LTD.	
23 March 2006 (23.03.2006)				
Applicant's or agent's file reference 27558		IMPORTANT NOTICE		
International application No. PCT/IL2004/000181	International filing date 24 February 20	e (day/month/year) F 04 (24.02.2004)	riority date (<i>day/month/year</i>) 27 April 2003 (27.04.2003)	
Applicant	PROTALIX	LTD. et al		
The International Bureau transmits herewith a c Treaty)			vility (Chapter I of the Patent Cooperation	
The International Bureau of W 34, chemin des Colombette 1211 Geneva 20, Switzerlar	s	Authorized officer Si	min Baharlou	

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PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (Chapter I of the Patent Cooperation Treaty)

(PCT Rule 44bis)

Applicant's or agent's file reference 27558	FOR FURTHER ACTION	See item 4 below	· · · · · · · · · · · · · · · · · · ·
International application No. PCT/IL2004/000181	International filing date (day/month/year) 24 February 2004 (24.02.2004)	Priority date (day/month/year) 27 April 2003 (27.04.2003)	
International Patent Classification (8th See relevant information in Form F	h edition unless older edition indicated) PCT/ISA/237		
Applicant PROTALIX LTD.			

1.	This international preliminary report on patentability (Chapter I) is issued by the International Bureau on behalf of the International Searching Authority under Rule 44 <i>bis</i> .1(a).					
2.	This REPORT consists of a total of 7 sheets, including this cover sheet.					
	In the attached sheets, any reference to the written opinion of the International Searching Authority should be read as a reference to the international preliminary report on patentability (Chapter I) instead.					
3.	. This report contains indications relating to the following items:					
	Box No. I	Basis of the report				
	Box No. II	Priority				
	Box No. III	Non-establishment of opinion with regard to novelty, inventive step and industrial applicability				
	Box No. IV	Lack of unity of invention				
	Box No. V	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
	Box No. VI	Certain documents cited				
	Box No. VII	Certain defects in the international application				
	Box No. VIII	Certain observations on the international application				
4.	The International Bureau will cont, except where the applicant date (Rule 44bis .2).	ommunicate this report to designated Offices in accordance with Rules 44 <i>bis</i> .3(c) and 93 <i>bis</i> .1 but makes an express request under Article 23(2), before the expiration of 30 months from the priority				

	Date of issuance of this report 13 March 2006 (13.03.2006)		
The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland	Authorized officer Simin Baharlou		
Facsimile No. +41 22 740 14 35	Telephone No. +41 22 338 71 30		

From the INTERNATIONAL SE		PATENT COOP		
To: G.E.EHRLICH G.E. EHRILICH (199		IORITY		PCT REC'D 20 FEE
11 MENACHEM BE RAMAT-GAN, ISRA	GIN STREET			VRITTEN OPINION OF THE TONAL SEARCHING AUTHORITY
				(PCT Rule 43bis.1)
·			Date of mailing (day/month/year	
Applicant's or agent's	file reference	10000	FOR FURTHE) 17 FEB 2006 RACTION See paragraph 2 below
27558 International applicati	on No.	International filing dat	e (day/month/year)	Priority date (day/month/year)
PCT/IL04/00181	i	-		
International Patent C	assification (IPC) o	24 February 2004 (24.) or both national classific:	ation and IPC	27 April 2003 (27.04.2003)
Applicant	12N 9700, 9714, 17 0.1, 325, 410,; 536	12, 1/20, 5/00, 15/00; C0 5/23.1, 23.4, 23.5; 800/29	07H 21/04; A01H 11/ 05	/00 and US Cl.: 435/4, 6, 41, 69.1, 183, 195,
METABOGAL, LTD				
1. This opinion cont.	ins indications rela	ating to the following ite		
, KZ		-		
Box No. I	Basis of the	opinion		
Box No. II	Priority			
Box No. II	l Non-establis	shment of opinion with r	egard to novelty, inv	entive step and industrial applicability
Вох №. Г		y of invention		
Box No. V	Reasoned sta applicability	atement under Rule 43 <i>bi</i> r; citations and explanation	s.1(a)(i) with regard ons supporting such a	to novelty, inventive step or industrial statement
Box No. V				
Box No. V	I Certain defe	ets in the international a	oplication	
Box No. V.		ervations on the internation	-	
2. FURTHER AC	rion			
Authority other the	in this one to be the	2 AULOODIV ("IPEA") P	EXCEPT that this doe: IPEA has notified to	l be considered to be a written opinion of the s not apply where the applicant chooses an he International Bureau under Rule $66.1bis(b)$ lered.
of Form PCT/ISA/	220 or before the ex	appropriate, with amend xpiration of 22 months f	imente hetora tha a	PEA, the applicant is invited to submit to the approximation of 3 months from the date of mailing whichever expires later.
For further options	see Form PCT/IS/	A/220.		
3. For further details,	see notes to Form I	PCT/ISA/220.		_
Name and mailing addr		Date of comple	tion of this opinion	Authorized officer
		····P.	of the second openant	TTILLA MA CLOO
Mail Stop PCT, Commissioner f P.O. Box 1450	pr Patents ginia 22313-1450	15 November 2	2005 (15.11.2005)	Manfulath N. Rao, Ph. D) Jok

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International application No.

PCT/IL04/00181

Box N	o. I Basis of this opinion				
1. With	regard to the language, this opinion has been established on the basis of:				
	the international application in the language in which it was filed				
	a translation of the international application into, which is the lange international search (Rules 12.3(a) and 23.1(b)).	uage of a translation furnished for the purposes of			
2. With inven	regard to any nucleotide and/or amino acid sequence disclosed in the int tion, this opinion has been established on the basis of:	ernational application and necessary to the claimed			
a.	type of material				
	a sequence listing				
	table(s) related to the sequence listing				
b.	format of material				
	on paper				
	in electronic form				
c.	time of filing/furnishing				
	contained in the international application as filed.				
	filed together with the international application in electronic form.				
	furnished subsequently to this Authority for the purposes of search.				
3.	In addition, in the case that more than one version or copy of a sequence lis or furnished, the required statements that the information in the subseque application as filed or does not go beyond the application as filed, as appro-	sting and/or table(s) relating thereto has been filed ent or additional copies is identical to that in the priate, were furnished.			
4. Additio	anal comments:				

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International application No.

Box N	No. IV Lack of unity of invention
1.	In response to the invitation (Form PCT/ISA/206) to pay additional fees the applicant has, within the applicable time lime paid additional fees paid additional fees under protest and, where applicable, the protest fee paid additional fees under protest but the applicable protest fee was not paid
	not paid additional fees
2.	This Authority found that the requirement of unity of invention is not complied with and chose not to invite the applicant pay additional fees.
3. Th	is Authority considers that the requirement of unity of invention in accordance with Rule 13.1, 13.2 and 13.3 is
	complied with
	not complied with for the following reasons: the lack of unity section of the International Search Report(Form PCT/ISA/210)
	any coordinate and maximutonial search Report(Form PC1/ISA/210)
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	· · ·
4. Consec	puently, this opinion has been established in respect of the following parts of the international application: all parts.
	the parts relating to claims Nos. <u>1-24.28-31,33-37 and 42</u>
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orm PCT/	ISA/237 (Box No. IV) (April 2005)

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International application No. PCT/IL04/00181

Ber N. Y. D.	· · ·			
Box No. V Reasoned statement under Rule 43 bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
1. Statement	······································			
Novelty (N)	Claims <u>4, 12-24, 42</u> Claims <u>1-3, 5-11, 28-31, 33-37</u>			
Inventive step (IS)	Claims <u>NONE</u> Claims <u>1-24, 28-31, 33-37, 42</u>			
Industrial applicability (IA)	Claims <u>1-24, 28-31, 33-37, 42</u> Claims <u>NONE</u>			
2. Citations and explanations: Please See Continuation Sheet				
orm PCT/ISA/237 (Box No. V) (April 2005)				

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International application No. PCT/IL04/00181

Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

V. 2. Citations and Explanations:

Claims 1-3, 5-11, 28-31, 33-37 lack novelty under PCT Article 33(2) as being anticipated by Martin et al. (DNA, 1988, Vol. 7, No.2, pages 99-106). Claims 1-3, 5-11, 28-31, 33-37 are drawn to a host cell producing a high mannose recombinant protein comprising a polynucleotide encoding the recombinant protein and a signal for causing the recombinant protein to be produced as a high mannose protein, wherein the polynucleotide comprises a first nucleic acid sequence encoding said protein of interest operably linked to a second nucleic acid sequence encoding a signal peptide wherein said signal peptide comprises a ER targeting peptide and wherein said host cell is a prokaryotic or a eukaryotic host cell and wherein said polypeptide is one of the lysosomal proteins such as glucocerebrosidase. Claim 28-31, 33-37 are also drawn to a recombinant biologically active high mannose lysosomal enzyme having at least one oligosaccharide chain comprising an exposed mannose residue. Martin et al. disclose one such host cell comprising a polynucleotide encoding said enzyme wherein said polypeptide is produced as a high-mannose protein in high levels. Matin et al. also disclose a recombinant glucocerebrosidase wherein said enzyme is inherently a biologically active high mannose lysosomal enzyme having at least one one oligosaccharide chain comprising an exposed mannose residue. Thus, Martin et al. anticipate claims 1-3, 5-11, 28-31, 33-37 as written.

Claims 4, 12-24 and 42 lack an inventive step under PCT Article 33(3) as being obvious over the prior art as applied in the immediately preceding paragraph and further in view of Boller et al. and Zhu et al. Claims 4, 12-24 and 42 are drawn to a host cell producing a high mannose recombinant protein comprising a polynucleotide encoding the recombinant protein and a signal for causing the recombinant protein to be produced as a high mannose protein, wherein the polynucleotide comprises a first nucleic acid sequence encoding said protein of interest operably linked to a second nucleic acid sequence with with SEQ ID NO:1 encoding a signal peptide wherein said signal peptide comprises a ER targeting peptide and wherein said polynucleotide is operably linked to a third polynucleotide sequence with SEQ ID NO:2 encoding a plant vacuolar targeting sequence, and wherein said host cell is a plant cell and wherein said polypeptide is one of the lysosomal proteins such as glucocerebrosidase. Claim 42 is drawn to a recombinant protein produced from a plant host cell. The reference of Martin et al. has already been discussed above. Martin et al. teach the production of glucocerebrosidase, a lysosomal protien recombinantly using a host cell comprising a polynucleotide with a signal sequence. The reference of Zhu et al. teach the polynucleotide encoding the signal peptide SEQ ID NO:1 and its use in producing novel recombinant proteins. On similar lines Boller et al. teach the vacuolar targeting sequence SEQ ID NO:2 and its use in targeting polypeptides into the vacuolar space. The invention as a whole is directed to production of glucocerebrosidase as a transgenic protein in plant host cells. The art and the above references teach and provide all sequences required for expressing the glucocerebrosidase as a transgenic protein. The production of mammalian proteins in plant products such as fruits and seed is well known since it eliminates the steps of purification and makes the recombinant protein ready for administration as a plant product. Therefore, with the above references in hand, it would have been obvious to one of ordinary skill in the art to produce human glucocerebrosidase, which is used in enzyme replacement therapy for lysosomal enzyme disorders, as a plant protein by expressing as a polynucleotide linked to the above signal sequence and vacuolar targeting sequences. One of ordinary skill in the art would have been motivated to do so since the lysosomal protein is extensively used in enzyme replacement therapy and production of the protein as a plant product would avoid the extensive purification steps and can be easily administered as a plant Form PCT/ISA/237 (Supplemental Box) (April 2005)

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International application No. PCT/IL04/00181

Supplemental Box In case the space in any of the preceding boxes is not sufficient. product. One of ordinary skill in the art would have had a reasonable expectation of success since Martin et al. already provide a host cell producing the high-mannose protein, Zhu et al. and Boller et al. provide the sequences to make a DNA construct to be expressed in a plant cell. Therefore the above invention would have been *prima facie* obvious to one of ordinary skill in the art. Claims 1-24, 28-31, 33-37, 42 meet the criteria set out in PCT Article 33(4), and thus have industrial applicability because the subject matter claimed can be made or used in industry.

Form PCT/ISA/237 (Supplemental Box) (April 2005)

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